## **CLAIMS**

1. A carbapenem compound represented by the following formula [1],

wherein  $R^1$  is  $C_1$ - $C_3$  alkyl group or  $C_1$ - $C_3$  alkyl group substituted by hydroxy group,

R is hydrogen atom or a group which reproduces carboxyl group by hydrolysis in vivo, and

G is a group represented by

the formula G1:

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the formula G2:

wherein Y¹ is C₁-C₄ alkyl, C₂-C₄ alkoxy, -(CH₂)<sub>ma</sub>-O-CH₃ (in which ma is an integer of 1~3), -O-(CH₂)<sub>ma</sub>-O-(CH₂)<sub>mb</sub>-CH₃ (in which ma is the same as defined above, mb is an integer of 0~3), trifluoromethoxy, halogen atom, cyano or -SO₂NR²R³ (in which R² and R³ are independently hydrogen atom, optionally substituted lower alkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aralkyl, or optionally substituted heteroarylalkyl, or R² and R³ may be taken together with the nitrogen atom to form a 3 to 7 membered hetero ring which may

be substituted.), or the formula G3:

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$$\begin{array}{c|c}
H & H \\
\hline
 & A \\
\hline
 & R^0
\end{array}$$
[G3]

wherein A is -(CH<sub>2</sub>)<sub>r</sub>-(in which r is an integer of 1~3), -(CH<sub>2</sub>)<sub>s</sub>-O-(CH<sub>2</sub>)<sub>t</sub>-( in which s and t are independently is an integer of 0~3), -O-(CH<sub>2</sub>)<sub>r</sub>-O-(CH<sub>2</sub>)<sub>s</sub>-(in which r and s are the same as defined above), -(CH<sub>2</sub>)<sub>s</sub>-NR<sup>a</sup>-(CH<sub>2</sub>)<sub>t</sub>-(in which, s and t are the same as defined above, Ra is hydrogen atom, protective group of amino group or optionally substituted C<sub>1</sub>-C<sub>6</sub> alkyl), Ro is hydrogen atom, the formula [2]:

$$\begin{array}{ccc}
O & R^{2a} \\
C N & R^{3a}
\end{array}$$
[2]

wherein R<sup>2a</sup> and R<sup>3a</sup> are independently (i) hydrogen atom, (ii) optionally substituted C<sub>1</sub>-C<sub>6</sub> alkyl, (iii) optionally substituted C<sub>3</sub>-C<sub>7</sub> cycloalkyl, (iv) optionally substituted aryl, (v) optionally substituted heteroaryl, (vi) optionally substituted aralkyl, (vii) optionally substituted heteroarylalkyl, or (viii) an optionally substituted 3 to 7 membered hetero ring, or R<sup>2a</sup> and R<sup>3a</sup> are taken together with the nitrogen atom to form a 3 to 7 membered hetero ring which may be substituted or the formula [3]:

$$\begin{array}{ccc}
O \\
II \\
C & (O)_{m}R^{3b}
\end{array}$$
 [3]

wherein m is an integer of 0 or 1,  $R^{3b}$  is hydrogen atom, optionally substituted  $C_1$ - $C_6$  alkyl, optionally substituted  $C_3$ - $C_7$  cycloalkyl, optionally substituted aryl, optionally substituted heteroaryl, optionally substituted aralkyl, optionally substituted heteroarylalkyl, or an optionally substituted 3 to 7 membered hetero ring, and when m is 1,  $R^{3b}$  may further mean a

group which reproduces carbonyl group by hydrolysis in vivo, provided that when t is 0 and m is 1, R<sup>3b</sup> is other group than hydrogen atom, and Y<sup>2</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, halogen atom, cyano or -NR<sup>4</sup>R<sup>5</sup> (in which R<sup>4</sup> and R<sup>5</sup> are independently

- (i) hydrogen atom, (ii) a protective group of amino group, (iii) optionally substituted C<sub>1</sub>-C<sub>6</sub> alkyl, (iv) optionally substituted C<sub>3</sub>-C<sub>7</sub> cycloalkyl, (v) formyl, (vi) C<sub>2</sub>-C<sub>7</sub> alkylcarbonyl, (vii) optionally substituted aryl, (viii) optionally substituted heteroaryl, (ix) optionally substituted aralkyl, (x) optionally substituted heteroarylalkyl, or (xi) an optionally substituted 3 to 7 membered hetero ring, or R<sup>4</sup> and R<sup>5</sup> are taken together with the nitrogen atom to form pyrrolidine, piperidine or azepam), or a pharmaceutically acceptable salt thereof.
- 2. A carbapenem compound or a pharmaceutically acceptable salt thereof represented by the following formula [1-a] wherein G is G1 in the above formula [1]:

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wherein R<sup>1</sup> and R are the same as defined n claim 1, or a pharmaceutically acceptable salt thereof.

3. A carbapenem compound represented by the following formula [1-b]:

$$R^1$$
 OMe [1-b]

wherein R1 and R are the same as defined in claim 1,

or a pharmaceutically acceptable salt thereof.

4. A carbapenem compound represented by the following formula [1-c]:

$$R^1$$
 [1-c]

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wherein  $R^1$ , R and  $Y^1$  are the same as defined in claim 1, or a pharmaceutically acceptable salt thereof.

5. A carbapenem compound or a pharmaceutically acceptable salt thereof represented by the following formula [1-d]:

$$\begin{array}{c|c}
R^1 & H & H \\
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wherein R<sup>1</sup>, R, A, R<sup>0</sup> and Y<sup>2</sup> are the same as defined in claim 1, or a pharmaceutically acceptable salt thereof.

6. The carbapenem compound claimed in any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof wherein a group which reproduces carboxyl group by hydrolysis in vivo is a group of the formula [4]:

$$\begin{array}{ccc}
\leftarrow \text{CH}_2\text{OC} & (O)_n & \mathbb{R}^7 \\
\downarrow & \downarrow \downarrow \\
\mathbb{R}^6 & O
\end{array}$$
[4]

wherein  $R^6$  is hydrogen atom or  $C_1$ - $C_6$  alkyl,  $R^7$  is optionally substituted  $C_1$ - $C_{10}$  alkyl, or optionally substituted  $C_3$ - $C_{10}$  cycloalky, and nis an integer of 0 or 1.

7. The carbapenem compound claimed in any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof wherein R is a group of the formula [4] claimed in claim 4.

- 8. The carbapenem compound claimed in any one of claims 1 to 7 or a pharmaceutically acceptable salt thereof wherein R<sup>1</sup> is 1-hydroxyethyl.
- 9. The carbapenem compound claimed in any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof wherein R is pivaloyloxymethyl, acetyloxymethyl, acetyloxy-1-ethyl, isopropyloxycarbonyloxy-1-ethyl or cyclohexyloxycarbonyloxy-1-ethyl.

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- 10. The carbapenem compound claimed in any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof wherein Ris pivaloyloxymethyl.
- 11. The carbapenem compound claimed in any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof wherein R is phthalidyl or (5-methyl-2-oxo-1,3-dioxol-4-yl)methyl.
- 12. The carbapenem compound claimed in any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof wherein R is hydrogen atom.
- 13. The carbapenem compound claimed in claim 4 or a pharmaceutically acceptable salt thereof wherein Y<sup>1</sup> is C<sub>2</sub>-C<sub>4</sub> alkoxy, (CH<sub>2</sub>)<sub>ma</sub>-O-CH<sub>3</sub> (in which ma is the same as defined in claim 1) or -O-(CH<sub>2</sub>)<sub>ma</sub>-O-(CH<sub>2</sub>)<sub>mb</sub>-CH<sub>3</sub> (in which ma and mb are the same as defined in claim 1).
- 14. The carbapenem compound claimed in claim 4 or a pharmaceutically acceptable salt thereof wherein Y<sup>1</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, trifluoromethoxy, halogen atom or cyano.
- 15. The carbapenem compound claimed in claim 4 or a pharmaceutically acceptable salt thereof wherein Y<sup>1</sup> is -SO<sub>2</sub>NR<sup>2</sup>R<sup>3</sup> (in which R<sup>2</sup> and R<sup>3</sup> are the same as defined in claim 1).
- 25 16. The carbapenem compound claimed in claim 4 or a pharmaceutically acceptable salt thereof wherein Y<sup>1</sup> is ethoxy, -CH<sub>2</sub>-O-CH<sub>3</sub>, -(CH<sub>2</sub>)<sub>2</sub>-O-CH<sub>3</sub> or -O-(CH<sub>2</sub>)<sub>2</sub>-O-CH<sub>3</sub>.
  - 17. The carbapenem compound claimed in any one of claims 4, 13 to 16 or a pharmaceutically acceptable salt thereof wherein Y<sup>1</sup> on benzene ring is metha or para to the binding position of 7-oxo-1-

azabicyclo[3.2.0]hept-2-ene.

- 18. The carbapenem compound claimed in any one of claims 4, 13 to 16 or a pharmaceutically acceptable salt thereof wherein Y<sup>1</sup> on benzene ring is para to the binding position of 7-oxo-1-azabicyclo[3.2.0]hept-2-ene.
- 19. The carbapenem compound claimed in claim 5 or a pharmaceutically acceptable salt thereof wherein R<sup>o</sup> is a formula [2]:

$$\begin{array}{ccc}
O & R^{2a} \\
CN & R^{3a}
\end{array}$$
[2]

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wherein R<sup>2a</sup> and R<sup>3a</sup> are the same as defined in claim 1.

20. The carbapenem compound claimed in claim 5 or a pharmaceutically acceptable salt thereof, wherein R<sup>o</sup> is a formula [3]:

wherein m and R3b are the same as defined in claim 1.

- 21. The carbapenem compound claimed in claim 5 or a pharmaceutically acceptable salt thereof wherein Y<sup>2</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl.
- 22. The carbapenem compound claimed in claim 5 or a pharmaceutically acceptable salt thereof wherein Y<sup>2</sup> is C<sub>1</sub>-C<sub>4</sub> alkoxy.
- 23. The carbapenem compound claimed in claim 5 or a pharmaceutically acceptable salt thereof wherein Y<sup>2</sup> is halogen atom or cyano.
- 24. The carbapenem compound claimed in claim 5 or a pharmaceutically acceptable salt thereof wherein Y<sup>2</sup> is -NR<sup>4</sup>R<sup>5</sup> (in which R<sup>4</sup> and R<sup>5</sup> are the same as defined in claim 1.
  - 25. A medicament containing a carbapenem compound claimed in any one of claims 1 to 24 or a pharmaceutically acceptable salt thereof as an active ingredient.
  - 26. An antibacterial agent containing a carbapenem compound claimed in any one of claims 1 to 24 or a pharmaceutically acceptable salt thereof as an active ingredient.

- 27. An oral medicament containing a carbapenem compound claimed in any one of claims 1 to 24 or a pharmaceutically acceptable salt thereof as an active ingredient.
- 28. An oral antibacterial agent containing a carbapenem compound claimed in any one of claims 1 to 24 or a pharmaceutically acceptable salt thereof as an active ingredient.

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